CLAIMS:

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- 1. A method of providing selective, substantially total, non-regenerative apoptosis of pancreatic acinar cells comprising a single-dose, subcutaneous or intra-arterial administration of a composition of cyanohydroxybutene and a pharmacologically acceptable aqueous carrier.
- 2.\A method according to claim 1, wherein said therapeutic window is selected to minimise liver damage in said patient.
- 3. A method according to claim 1 or 2, wherein said administration is subcutaneous.
- 4. A method according to any one of claims 1 and 3, wherein said cyanohydroxybutene is administered at a dosage within the range of 140-160 mg

 CHB/kg of body weight.

on the basis of said pancreatic acinar cells including acinar carcinoma cells.

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6. A method for treating pancreatic disease including administering to a patient a single-dose, subcutaneous or intra-arterial, therapeutically effective amount of cyanohydroxybutene wherein said amount is sufficient to cause selective, substantially total, substantially non-regenerative apoptosis of acinar cells in the patient.



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8. A method as claimed in claim 7 wherein the CHB dose is within a range of 125-/160 mg CHB/kg of body weight.

said subject in an amount sufficient to cause selective, substantially total,

substantially non-regenerative apoptosis of malignant acinar cells in a patient.

- 9. A method as claimed in claim 8 wherein the CHB dose is within the range of 140-160 mg CHB/kg of body weight.
- 10. A method as claimed in claim 7 wherein the carcinoma involves either acinar cell carcinoma or pancreatic carcinoma containing a mixed population of cells including acinar cells.
- 11. A method as claimed in claim 7 wherein said CHB molecule is conjugated to a ligand which is selected to to bind to an acinar cell surface receptor.
 - A method according to any one of claims 7 to 11, wherein said dose is selected whereby liver damage in the subject is minimised.
- 13. A method of treating acute or chronic pancreatitis including the steps of: preparing a cyanohydroxybutene (CHB) formulation; and

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AMENDED SHEET

administering a subcutaneous or intra-arterial single dose of a CHB formulation to said subject in an amount sufficient to cause selective, substantially total, substantially non-regenerative apoptosis of malignant acinar cells in a patient.

- 14. A method of treating acute or chronic pancreatitis as claimed in claim 13 wherein the CHB dose is within a range of 125-160 mg CHB/kg of body weight.
- 15. A method of treating acute or chronic pancreatitis as claimed in claim 13 or 14 wherein the CHB formulation is administered by subcutaneous injection.

A method according to any one of claims 13 to 15, wherein said dose is selected whereby liver damage in the subject is minimised.

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